



## Complete Summary

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### **GUIDELINE TITLE**

HIV screening in pregnancy.

### **BIBLIOGRAPHIC SOURCE(S)**

Keenan-Lindsay L, Yudin MH, Boucher M, Cohen HR, Gruslin A, MacKinnon CJ, Money DM, Paquet C, Steben M, van Schalkwyk J, Wong T, Maternal Fetal Medicine Committee, Society of Obstetricians and Gynaecologists of Canada. HIV screening in pregnancy. J Obstet Gynaecol Can 2006 Dec;28(12):1103-7. [35 references] [PubMed](#)

### **GUIDELINE STATUS**

This is the current release of the guideline.

## COMPLETE SUMMARY CONTENT

SCOPE  
METHODOLOGY - including Rating Scheme and Cost Analysis  
RECOMMENDATIONS  
EVIDENCE SUPPORTING THE RECOMMENDATIONS  
BENEFITS/HARMS OF IMPLEMENTING THE GUIDELINE RECOMMENDATIONS  
QUALIFYING STATEMENTS  
IMPLEMENTATION OF THE GUIDELINE  
INSTITUTE OF MEDICINE (IOM) NATIONAL HEALTHCARE QUALITY REPORT  
CATEGORIES  
IDENTIFYING INFORMATION AND AVAILABILITY  
DISCLAIMER

## SCOPE

### **DISEASE/CONDITION(S)**

Human immunodeficiency virus (HIV) infection in pregnancy

### **GUIDELINE CATEGORY**

Counseling  
Evaluation  
Prevention  
Risk Assessment  
Screening

### **CLINICAL SPECIALTY**

Infectious Diseases  
Internal Medicine  
Obstetrics and Gynecology  
Pediatrics  
Preventive Medicine

## **INTENDED USERS**

Health Care Providers  
Patients  
Physicians  
Public Health Departments

## **GUIDELINE OBJECTIVE(S)**

- To provide recommendations to obstetric health care providers and to minimize practice variations for human immunodeficiency virus (HIV) screening, while taking provincial and territorial recommendations into account
- To provide health care providers and pregnant women with information about HIV screening in pregnancy

## **TARGET POPULATION**

Pregnant women

## **INTERVENTIONS AND PRACTICES CONSIDERED**

1. Human immunodeficiency virus (HIV) screening
2. Individualized counseling if indicated
3. HIV prophylaxis for women at high risk and unknown status
4. Postpartum infant prophylaxis if indicated
5. Follow-up of HIV-positive women

## **MAJOR OUTCOMES CONSIDERED**

- Vertical transmission of human immunodeficiency virus (HIV) infection rate
- Seroprevalence rate
- HIV screening rate

## **METHODOLOGY**

### **METHODS USED TO COLLECT/SELECT EVIDENCE**

Searches of Electronic Databases

### **DESCRIPTION OF METHODS USED TO COLLECT/SELECT THE EVIDENCE**

The Cochrane Library and Medline were searched for English-language articles published related to human immunodeficiency virus (HIV) screening and

pregnancy. Additional articles were identified through the references of these articles. All study types were reviewed.

## **NUMBER OF SOURCE DOCUMENTS**

Not stated

## **METHODS USED TO ASSESS THE QUALITY AND STRENGTH OF THE EVIDENCE**

Weighting According to a Rating Scheme (Scheme Given)

## **RATING SCHEME FOR THE STRENGTH OF THE EVIDENCE**

### **Quality of Evidence Assessment\***

**I:** Evidence obtained from at least one properly randomized controlled trial

**II-1:** Evidence from well-designed controlled trials without randomization

**II-2:** Evidence from well-designed cohort (prospective or retrospective) or case-control studies, preferably from more than one centre or research group

**II-3:** Evidence obtained from comparisons between times or places with or without the intervention. Dramatic results in uncontrolled experiments (such as the results of treatment with penicillin in the 1940s) could also be included in this category

**III:** Opinions of respected authorities, based on clinical experience, descriptive studies, or reports of expert committees

\*Adapted from the Evaluation of Evidence criteria described in the Canadian Task Force on Periodic Health Exam.

## **METHODS USED TO ANALYZE THE EVIDENCE**

Systematic Review

## **DESCRIPTION OF THE METHODS USED TO ANALYZE THE EVIDENCE**

Not stated

## **METHODS USED TO FORMULATE THE RECOMMENDATIONS**

Expert Consensus

## **DESCRIPTION OF METHODS USED TO FORMULATE THE RECOMMENDATIONS**

Not stated

## RATING SCHEME FOR THE STRENGTH OF THE RECOMMENDATIONS

### Classification of Recommendations\*

- A. There is good evidence to support the recommendation that the condition be specifically considered in a periodic health examination.
- B. There is fair evidence to support the recommendation that the condition be specifically considered in a periodic health examination.
- C. There is poor evidence regarding the inclusion or exclusion of the condition in a periodic health examination.
- D. There is fair evidence to support the recommendation that the condition not be considered in a periodic health examination.
- E. There is good evidence to support the recommendation that the condition be excluded from consideration in a periodic health examination.

\*Adapted from the Evaluation of Evidence criteria described in the Canadian Task Force on Periodic Health Exam.

## COST ANALYSIS

Guideline developers reviewed published cost analyses.

## METHOD OF GUIDELINE VALIDATION

Internal Peer Review

## DESCRIPTION OF METHOD OF GUIDELINE VALIDATION

This guideline has been reviewed by the Maternal Fetal Medicine Committee and approved by the Executive and Council of the Society of Obstetricians and Gynaecologists of Canada.

## RECOMMENDATIONS

### MAJOR RECOMMENDATIONS

Definitions of the levels of evidence (I, II-1, II-2, II-3, and III) and grades of recommendations (A-E) are provided at the end of the "Major Recommendations" field.

#### Counselling

1. All pregnant women should have human immunodeficiency virus (HIV) screening with appropriate counselling. This testing must be voluntary. Screening should be considered a standard of care, although women must be informed of the policy, its risks and benefits, and the right of refusal. Women must not be tested without their knowledge. **(II-2 B)**
2. Pre-test counselling and the patient's decision about testing should be documented in the patient's chart. **(III-B)**
3. Women who decline screening should still have concerns discussed and should continue to receive optimum antenatal care. **(III-C)**

## **When to Provide Screening**

4. Women should be offered HIV screening at their first prenatal visit. (**I-A**)
5. Women who test negative for HIV and continue to engage in high-risk behaviour should be retested in each trimester of pregnancy. (**II-3 B**)
6. Women with no prenatal care and unknown HIV status should be offered testing when admitted to hospital for labour and delivery. Women at high risk for HIV and with unknown status should be offered HIV prophylaxis in labour, and HIV prophylaxis should be given to the infant post partum. (**III-B**)

## **Appropriate Follow-Up**

7. Women who test positive for HIV should be followed by practitioners who are knowledgeable in the care of HIV-positive women. (**III-C**)

## **Definitions:**

### **Quality of Evidence Assessment\***

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**III:** Opinions of respected authorities, based on clinical experience, descriptive studies, or reports of expert committees

### **Classification of Recommendations\***

- A. There is good evidence to support the recommendation that the condition be specifically considered in a periodic health examination.
- B. There is fair evidence to support the recommendation that the condition be specifically considered in a periodic health examination.
- C. There is poor evidence regarding the inclusion or exclusion of the condition in a periodic health examination.
- D. There is fair evidence to support the recommendation that the condition not be considered in a periodic health examination.
- E. There is good evidence to support the recommendation that the condition be excluded from consideration in a periodic health examination.

\*The quality of evidence reported in these guidelines has been adapted from the Evaluation of Evidence criteria described in the Canadian Task Force on the Periodic Health Exam.

\*\*Recommendations included in these guidelines have been adapted from the Classification of Recommendations criteria described in the Canadian Task Force on the Periodic Health Exam.

## **CLINICAL ALGORITHM(S)**

None provided

## **EVIDENCE SUPPORTING THE RECOMMENDATIONS**

### **TYPE OF EVIDENCE SUPPORTING THE RECOMMENDATIONS**

The type of supporting evidence is identified and graded for each recommendation (see "Major Recommendations").

## **BENEFITS/HARMS OF IMPLEMENTING THE GUIDELINE RECOMMENDATIONS**

### **POTENTIAL BENEFITS**

- Decrease in the rate of vertical transmission of human immunodeficiency virus (HIV) from mother to fetus
- Confirmation of HIV infection in the woman, which allows optimization of her health and long-term management

### **POTENTIAL HARMS**

Not stated

## **QUALIFYING STATEMENTS**

### **QUALIFYING STATEMENTS**

This guideline reflects emerging clinical and scientific advances as of the date issued and is subject to change. The information should not be construed as dictating an exclusive course of treatment or procedure to be followed. Local institutions can dictate amendments to these opinions. They should be well documented if modified at the local level.

## **IMPLEMENTATION OF THE GUIDELINE**

### **DESCRIPTION OF IMPLEMENTATION STRATEGY**

An implementation strategy was not provided.

## **INSTITUTE OF MEDICINE (IOM) NATIONAL HEALTHCARE QUALITY REPORT CATEGORIES**

### **IOM CARE NEED**

Living with Illness  
Staying Healthy

## **IOM DOMAIN**

Effectiveness  
Patient-centeredness

## **IDENTIFYING INFORMATION AND AVAILABILITY**

### **BIBLIOGRAPHIC SOURCE(S)**

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### **ADAPTATION**

Not applicable: The guideline was not adapted from another source.

### **DATE RELEASED**

2006 Dec

### **GUIDELINE DEVELOPER(S)**

Society of Obstetricians and Gynaecologists of Canada - Medical Specialty Society

### **SOURCE(S) OF FUNDING**

Society of Obstetricians and Gynaecologists of Canada

### **GUIDELINE COMMITTEE**

Infectious Disease Committee

### **COMPOSITION OF GROUP THAT AUTHORED THE GUIDELINE**

*Principal Authors:* Lisa Keenan-Lindsay, RN, MN, Toronto ON; Mark H. Yudin, MD, MSc, FRCSC, Toronto ON

*Committee Members:* Marc Boucher, MD, FRCSC, Montreal QC; Howard Ronald Cohen, MD, FRCSC, Toronto ON; Andrée Gruslin, MD, FRCSC, Ottawa ON; Catherine Jane MacKinnon, MD, FRCSC, Brantford ON; Deborah M. Money, MD, FRCSC, Vancouver BC; Caroline Paquet, RM, MSc, Trois-Rivières QC; Marc Steben, MD, Montreal QC; Julie van Schalkwyk, MD, FRCSC, Vancouver BC; Thomas Wong, MD, MPH, FRCPC, Ottawa ON; Mark H. Yudin, MD, MSc, FRCSC, Toronto ON

## **FINANCIAL DISCLOSURES/CONFLICTS OF INTEREST**

Not stated

## **GUIDELINE STATUS**

This is the current release of the guideline.

## **GUIDELINE AVAILABILITY**

Electronic copies: Available in Portable Document Format (PDF) from the [Society of Obstetricians and Gynaecologists of Canada Web site](#).

Print copies: Available from the Society of Obstetricians and Gynaecologists of Canada, La société des obstétriciens et gynécologues du Canada (SOGC) 780 promenade Echo Drive Ottawa, ON K1S 5R7 (Canada); Phone: 1-800-561-2416

## **AVAILABILITY OF COMPANION DOCUMENTS**

None available

## **PATIENT RESOURCES**

None available

## **NGC STATUS**

This NGC summary was completed by ECRI Institute on February 10, 2009. The information was verified by the guideline developer on March 4, 2009.

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